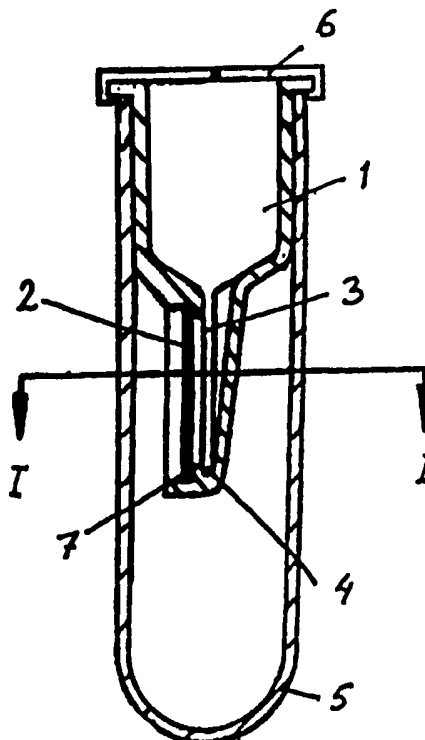




## INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

<p>(51) International Patent Classification<sup>5</sup>: B01L 3/14, B01D 61/18, 63/16, 29/05, 29/74</p>	<p>A1</p>	<p>(11) International Publication Number: WO 94/27724 (43) International Publication Date: 8 December 1994 (08.12.94)</p>
<p>(21) International Application Number: PCT/EP94/01701 (22) International Filing Date: 24 May 1994 (24.05.94) (30) Priority Data: 9301759-8 21 May 1993 (21.05.93) SE (71)(72) Applicant and Inventor: VASSAROTTI, Vincenzo [IT/CH]; Au Village, CH-1180 Bugnax sur Rolle (CH). (74) Agent: JOHANSSON, Lars; Patech S.A., Le Haut-des-Champs, CH-1173 Fechy (CH).</p>		<p>(81) Designated States: JP, US, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE).  Published With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</p>
<p>(54) Title: CENTRIFUGAL METHOD FOR CONCENTRATING MACROMOLECULES FROM A SOLUTION AND DEVICE FOR CARRYING OUT SAID METHOD</p> <p>(57) Abstract</p> <p>Method and device for concentrating macromolecules from a solution by means of spinning a concentrator device in a centrifuge. Said device comprises a sample reservoir (1) for receiving a sample to concentrate or purify, filter means for filtering the sample, a centrifuge tube (5) for collecting the filtrate and a concentrate pocket (4) for retention of concentrate. Typically the sample reservoir (1) of the device is provided with a portion constituting a filtration chamber (3) with a volume being only a small fraction of the total volume of the sample reservoir and the filter means is arranged in the sidewall and the concentrate pocket (4) is arranged in the bottom wall of said filtration chamber. In the method the liquid sample is put into the sample reservoir and the unit comprising the sample reservoir (1) and the filter means (2) is put at least partly into the centrifuge tube for collecting the filtrate. This assembly is centrifuged whereby a force vector is created acting on the macromolecules and sweeping the filter surface, the macromolecules are collected in a concentrate pocket (4) away from the filter surface thereby eliminating the risk of concentration to dryness.</p> 		

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CENTRIFUGAL METHOD FOR CONCENTRATING MACROMOLECULES FROM A  
SOLUTION AND DEVICE FOR CARRYING OUT SAID METHOD

- 5 This invention relates to centrifugal methods and devices for  
concentrating macromolecules from a solution.

There has been a number of analytical procedures developed in  
the biochemical art wherein it is required to remove solvent  
10 from e.g. protein solutions in order to have a more  
concentrated protein sample which can be analysed  
effectively, or in order to have a protein free filtrate for  
analysis, or to replace or remove low molecular weight ions  
or solutes, or to study protein binding attributes of various  
15 chemicals in combination with various protein samples. Many  
other analytical procedures involving not only proteins but  
macromolecular species in general, have also been developed  
wherein it is necessary to concentrate a macromolecular  
component in a liquid sample.

20

When concentrating small volumes of macromolecules in  
solution using filtration, e.g. ultrafiltration, there exist  
the problems of macromolecular binding to the membrane  
surface and of filtration to dryness. If part of the  
25 retentate is adsorbed on the membrane surface the yield of  
the process will of course decrease and the filtration time  
will increase due to the decreasing effective membrane area.

In the design of centrifugal filtration devices, the  
30 effective membrane area to sample volume to be filtered has  
been limited by the designs proposed. Low membrane area to  
volume ratios provide good recovery but low filtration rates.  
Conversely low volume and large membrane surfaces result in  
faster filtration but lower recoveries due to the protein  
35 binding to the membrane surface.

In the prior art, the centrifugal self-cleaning ultrafilter, described in US-A-3 488 768, the teachings of which are incorporated herein, is limited due to the large membrane surface area which causes excessive adsorptive loss of the macromolecules.

The initial advantage of a high flow rate through a large filter surface area is rapidly limited as the concentration goes on and the physical surface area is decreasing. The flow rate through the remaining filter area is also reduced by the effect of concentration polarization.

On the other hand, more recent designs such as AMICON MPS-1 Micropartition system (AMICON Publication 472) and the AMICON CENTRICON device described in US-A-4 632 761 limits membrane surface area to the area at the bottom of the sample reservoir tube, frequently resulting in a poor ratio of membrane surface area to sample volume and extended concentration time.

The dead stop feature which stops the sample concentrating to dryness according to US-A-4 632 761 is limited by remaining adsorptive losses which result from the concentrated sample remaining on the surface of the membrane and a reduction of effective membrane surface area resulting from this feature.

To recover the retentate from the concentrate pocket a reverse spinning step is used which means that the centrifuge has to be stopped and the sample reservoir with the filtering means and the centrifuge tube have to be taken out from the centrifuge. The different devices are then rearranged and again inserted into the centrifuge now in a way allowing the centrifugal forces to recover the retentate from the pocket at the bottom of the sample reservoir and the surface of the

filter. The use of a pipette to recover the retentate from the bottom of the sample reservoir is delicate because the surface of the filter might be damaged by the tip of the pipette. This necessary step to get the final product out of the concentration device is thus complicated and will take a lot of time. Furthermore, a dedicated retentate recovery tube has to be used which increases the costs.

Whilst the reverse spinning feature partially resolves the problem of loss of recovery on the membrane surface, subsequent loss is caused by the adsorption of the concentrated solution on the membrane and reservoir walls of the apparatus during the reverse spinning procedure.

In addition this device requires a specifically designed filtrate collection vessel which is mounted below the sample reservoir - filter assembly. Typical laboratory centrifuge tubes may not be used which adds costs for using this device.

It is an object of the invention to provide a centrifugal recovery process for the recovery of a maximal amount of concentrated macromolecular retentate without the need for an additional reverse spinning step to free residual particulate matter from the membrane surface and the retentive reservoir.

It is another object of the invention to provide a centrifugal recovery process compatible with a broad range of standard laboratory centrifuge tubes eliminating the need for product specific filtrate vessels.

It is another object of the invention to provide a process which allows the use of both lower cost swinging bucket centrifuge rotors as well as fixed angle versions providing increased flexibility in use.

It is a further object of this invention to provide a centrifugal concentrator which offers a high degree of flexibility in optimizing the relationship between the filtration rate and the decrease of macromolecular recovery  
5 caused by large membrane surfaces.

It is another object of the invention to provide a concentrator having a low volume filtration chamber preferably in the form of a thin channel, in the side wall of  
10 which the membrane is mounted. The channel being arranged below the main part of the sample reservoir which maximizes sample contact time with the total available membrane surface during concentration.

15 It is another object of the invention to provide a method which drives particulate matter away from the membrane surface to the concentrate pocket at the bottom of the filtration chamber at an early stage of centrifugation.

20 It is another object of this invention to provide a concentrator which gives constant final retentate volume in a separate concentrate pocket away from the membrane surface.

It is a still further object of this invention to provide a  
25 centrifugal micro volume concentrator that is economical to manufacture.

The problems of the prior art are overcome by the provision of a centrifugal method and a device for carrying out said  
30 method as defined in the appended claims.

In one embodiment of the centrifugal filtration system the membrane employed is an ultrafiltration or microfiltration membrane having a maximum pore size of one micron. The  
35 membrane could be in sheet form mounted on one or more sides

of the filtration chamber. In another embodiment, the membrane is tubular in shape and performs as an integral filtration chamber mounted below the sample reservoir. The membrane is mounted parallel to or within a maximum angle of 35 degrees with respect to the axis of the centrifuge tube. The filtration system could further comprise a thin channel of a maximum of 5 millimeters in the area of the filtration chamber to maintain a constant active membrane surface throughout most of the concentration procedure. This channel is preferably provided with a wider section to facilitate entry with a pipette for concentrate removal. The sample reservoir could be provided with a vented cap to reduce sample evaporation whilst preventing air locking during centrifugation.

15

In the method of the invention for concentrating macromolecules from a solution without excessive adsorption on the membrane surface and without filtering to dryness, the sample reservoir of the above described novel microconcentrator device is preferably placed in a swinging bucket centrifuge rotor so that during centrifugation the centrifugal force vector sweeps the membrane at an angle in the interval  $0 - 35^\circ$  and a concentrated macromolecular final retentate volume is obtained by centrifuging the microconcentrator device until the retentate level falls below the level of the membrane. A retentate reservoir below the membrane surface retains the final retentate solution eliminating the risk of concentrating to dryness.

30 If an embodiment of the concentrating device with the filter arranged in a window in the side wall of the filtration chamber is used with a fixed angle centrifuge certain combinations of on one hand the angle between the filter and the axis of the centrifuge tube and on the other hand the angle of the centrifuge rotor are especially interesting.

Combinations could be chosen so that the centrifugal force vector sweeps the membrane at an angle of approximately  $0^\circ$ , i.e. parallel to the filter surface, or in a direction away from the filter surface.

5

The retentate obtained by the method for concentration of macromolecules is preferably recovered by drawing the retentate from the concentrate reservoir by means of a pipette or other recovery device. No reverse spinning step is  
10 thus necessary in the method according to the invention.

Other characteristics and advantages of the invention will be apparent from the following description of a few embodiments which will be given with reference to the drawing on which,

15

FIG. 1 is a schematic cross section view of the microconcentrator device fitted into a filtrate recovery tube.

20 FIG. 2 is a sectional view of the microconcentrator device at the level I-I according to FIG. 1

FIG. 3 is a schematic view of the microconcentrator device showing the membrane support frame.

25

FIG. 4 is a schematic cross section of part of the membrane, membrane support and the joint fixed in the side wall of the concentrator device.

30 FIG. 5 is a schematic view in partial cross-section of a microconcentration device, using a tubular membrane and membrane support, fitted in a filtrate recovery tube

FIG. 6 is a schematic view in partial cross-section of a  
35 microconcentration device showing the principle of cascading

filter means of the same type as in figure 5.

Fig. 7 shows a further embodiment of the invention with a tapered sleeve on the outside of the filtration chamber.

5

Fig. 8 shows in a sectional view more in detail the essential parts of the embodiment according to fig. 7.

Referring to FIG. 1 there is shown an embodiment of the  
10 microconcentrator device of the invention comprising a sample  
reservoir 1, a membrane and support assembly 2 arranged in  
the side wall of a filtration chamber 3, which at the lower  
portion is provided with a retentate reservoir 4 for  
collecting the concentrate. The assembly is placed in a  
15 filtrate recovery tube 5 and a vented cap 6 closes the tube  
5. The retentate reservoir is preferably integrated into the  
design of the filtration chamber 3 and its uppermost edge  
aligns with the lower edge of the membrane window 7 in such a  
manner that when filtration is arrested by lack of useful  
20 membrane surface the retentate reservoir is full.

Preferably this reservoir is conical in shape to facilitate  
complete retentate removal and can be varied in size to best  
suit the concentration level required for different  
25 applications. Preferably the retentate reservoir 4 will be  
sized to allow concentrations of 10 to 300 times of the  
initial sample volume. The angle between the axis of the  
sample reservoir, coinciding with the axis of the centrifugal  
tube, and the flat membrane in the side wall of the  
30 filtration chamber can be chosen in the approximate interval  
0 - 35°.

A big advantage with the device according to the invention is  
that standard laboratory centrifuge tubes may be used as  
35 filtrate collection vessels. The sample reservoir including



the filtration chamber may be designed to be put on top of, partly into or completely into the filtrate collection vessel. Figure 1 shows the embodiment with the sample reservoir and the filtration chamber completely inside the  
5 filtrate collection vessel.

FIG. 2 shows a cross section of the filtration chamber with a pipette entry port 8, a channelled membrane support plate 21, the membrane 22, a porous sinter membrane support 23, and a  
10 membrane gasket 24. The filtration chamber 3 in the form of a flat channel is preferably of a thin section of less than 5 mm to maintain a high membrane surface to volume ratio as the concentration progresses.

15 If we define a concentration factor as the ratio between the initial sample volume and the final retentate volume the method and device according to the invention can reach much higher values of this factor than the methods and devices according to the prior art. This is due to the feature with a  
20 thin channel as filtration chamber and the retentate reservoir away from the filter surface at the bottom of the filtration chamber.

The entry port 8, allows for the removal of the concentrated  
25 retentate by means of a pipette. The channelled membrane support plate 21, is sealed to the sample reservoir 1, compressing the gasket 4 against the membrane and thereby providing a liquid tight seal. Any means of sealing may be used such as bonding the membrane directly to the membrane  
30 support frame, and thereby eliminating the need for the gasket 4. Similarly, the membrane support plate 23, may be sealed to the sample reservoir 1, by alternative means such as adhesive, heat or chemical bonding, insert moulding and interference fit.

FIG. 3 shows a schematic side view of the device, with the sample reservoir 1, the membrane window 7, and the filtrate outlet channels 9. The window provides for the fitting of the gasket 24, and the membrane 22. The filtrate outlet channels 5 9, provide for the evacuation of the filtrate from the membrane support to the filtrate recovery tube 5. The membrane window 7, can be sized, depending on the characteristics of the process solution and filtration rate of the membrane to achieve the best compromise between high 10 filtration speed and low adsorption on the membrane surface. When very high filtration rates are necessary, one or more additional membrane windows may be arranged on the other parts of the side wall of the filtration chamber.

15 FIG. 4, shows a detail of the membrane assembly with the sample reservoir 1, the membrane support plate 21 with incorporated filtrate outlet channels 9, the gasket seal 24, the membrane 22 and a porous sinter membrane support 23. The porous sinter membrane support 23 is preferably used to 20 provide a more homogeneous support for fragile membranes that could distort into the filtrate outlet channels of the membrane support plate. Any system of filtrate outlet channels may be employed as long as sufficient support is maintained for the membrane and at least one filtrate passage 25 is available to allow filtrate flow from the membrane to the filtrate reservoir 5.

In an advantageous further embodiment of the invention, cf. figures 7 and 8, the sample reservoir 1 and the filtration 30 chamber 3 including the retentate reservoir 4 form an integral part. The external form of the filtration chamber 3 could e.g. be half a cylinder with the flat surface constituting the membrane window 7. In the illustrations of figures 7 and 8 the actual form of the filtration chamber is 35 tapered in the direction of the retentate reservoir (cf.

below). The membrane 22 being fixed by suitable means against the window. On the outside of the filtration chamber 3 a corresponding half cylindrical sleeve 25 is arranged keeping the membrane firmly fixed against the window. The flat side  
5 26 of this half cylindrical sleeve has the role of the membrane support plate 21 in the embodiment described above. This means that the internal flat surface 27 of the sleeve is provided with the filtrate outlet channels 9 in the form of vertical grooves. The vertical position of the sleeve  
10 relative to the filtration chamber can be secured by means of cooperating snap-in arrangements (not shown) in the side wall of the chamber and the sleeve.

In a further development of this embodiment the external form  
15 of the filtration chamber 3 could be conical or more generally have a tapered form in the direction of the retentate reservoir, as actually shown in the figures 7 and 8. An angle of about  $3^\circ$  for the tapering has been giving excellent results. With a corresponding internal form of the  
20 sleeve 25 a very good fixation of the membrane will be possible. It has been shown that the membrane as such does not even have to be fixed against the window by means of glue, heat or the equivalent but will be kept in place tightly against the window by means of the forces exerted by  
25 the sleeve when this is slipped over the filtration chamber.

In figure 8 another feature of the invention is also shown. For the application with a swinging-bucket rotor the bottom of the reservoir has been arranged at an angle of about  $5^\circ$   
30 to the horizontal, which makes the last part of the volume of the liquid slide down into the filtration chamber or channel.

To allow flow extraction with fixed angle rotors with an angle up to  $45^\circ$  the corresponding angle at the bottom of the  
35 sample reservoir would be arranged at about  $46^\circ$

It could sometimes be of interest during the concentration process to check the volume achieved. The fact that the filtration chamber in the device according to the invention, especially according to figures 1 and 7, has the form of a thin channel containing a very small volume, sometimes in the order of 50 microliters, makes it possible to get an accurate reading of the remaining volume by putting indication markings on the sidewall of the channel or on the outside or inside of the sleeve. It is here assumed that the sleeve as well as the filtration chamber are made of transparent material.

The dead stop feature is realised by means of a partition wall 28 separating the retentate reservoir 4 from the surface of the membrane 22. It could be of advantage to give the cross section of the upper part of this wall the shape of a ramp as shown in figure 8 or a trapezoid.

FIG. 5 shows a schematic cross-section of an alternative design for the microfiltration device with the sample reservoir 1', the filtrate recovery tube 5', a tubular membrane 10 supported by means of a porous cylindrical tube 11 forming the side wall of the filtration chamber, a membrane sealing unit 12, with integral retentate reservoir 13 and two sealing gaskets 14, one for each end of the cylindrical tube. In this design the membrane sealing unit is screwed onto the sample reservoir 1' and tightens the sealing gaskets to provide a liquid tight seal between the membrane 10, the sample reservoir 1' and the membrane sealing unit 12. Any other means of sealing may be used including adhesive, heat or chemical bonding or a bayonet assembly. The tubular membrane 10 and its support may be increased or decreased in both diameter and length to achieve the best compromise between a fast filtration speed and minimum adsorption on the

membrane surface.

In a further embodiment of the invention, shown in figure 6, two or more microconcentrator devices may be stacked in a suitable filtrate recovery tube to sequentially fractionate a sample solution into two or more fractions containing macromolecules of different size. The figure shows a device based on the embodiment according to figure 5 with the cylindrical filtration chamber but other types of filtration chambers can alternatively be used. Length and diameter of the filters, i.e filter area, as well as the volume of the retentate reservoir can of course be chosen to suit the actual application. Preferably the successive filter units are screwed on top of each other and sealed from each other so that filtrate will not leak from one reservoir to the other during the handling. In the figure the thread for fastening the second device to the first one is not shown. The sealing is carried out by means of an O-ring 30 but any other appropriate sealing would do. In this process, the uppermost device will be fitted with a more open filter that retains the larger macromolecules but allows passage of smaller macromolecules that will be retained by a filter of lower porosity in the lower device. Depending on the application, up to five or even more devices may be stacked together with sequentially tighter filters. Due to the fact that in fixed angle rotors as well as in swinging bucket rotors the "lower" filters will be spinning at a greater distance from the spinning shaft of the centrifugator the centrifugal forces will be greater in these filters and compensate for the lower flow rate due to the tighter filters.

The fractionated samples may then be collected by means of a pipette individually from the retentate reservoir in each device.

- Although any type of semi permeable membrane may be used in the microconcentrator device of the invention, anisotropic membranes, i.e. membranes having an extremely thin
- 5 microporous barrier layer and a relatively thick macroporous support layer, are preferred and microporous or ultrafiltration membranes characterised by a maximum pore size of one micron has been shown to be most suitable.
- 10 The microconcentrator according to the invention is preferably used in a swinging-bucket centrifuge rotor, but the device may also be used in a fixed angle centrifuge rotor. When used in a swinging-bucket rotor the angle between the membrane surface and the direction of the force on the
- 15 molecules due to the spinning will be approximately  $0^\circ$ , which gives a very good cleaning effect on the membrane surface.

The most common fixed angle rotors use angles of 26-55 degrees. When used in these rotors the angle between the

20 membrane surface of the microconcentrator device and the force vector in the direction away from the membrane will reside in the interval between approximately 35 and 64 degrees which efficiently will keep especially the heavy molecules away from the membrane surface.

- 25 Other angles of the fixed rotor could of course be envisaged as well as other angles between the axis of the sample reservoir and the membrane surface in the side wall of the filtration chamber which will result in other sweeping angles
- 30 for the force vector without leaving the idea of the present invention.

## CLAIMS

1. Method for concentrating macromolecules from a solution  
5 by means of spinning a concentrator device in a centrifuge,  
said device comprising a sample reservoir (1) for receiving a  
sample to concentrate or purify, filter means for filtering  
the sample, a centrifuge tube (5) for collecting the filtrate  
and a concentrate pocket (4) for retention of concentrate, in  
10 which method the liquid sample is put into the sample  
reservoir characterised in that it additionally comprises the  
following steps: the unit comprising the sample reservoir (1)  
and the filter means (2) is put at least partly into the  
centrifuge tube for collecting the filtrate, the thus created  
15 assembly is centrifuged whereby a force vector is created  
acting on the macromolecules and sweeping the filter surface,  
the macromolecules are collected in a concentrate pocket (4)  
away from the filter surface thereby eliminating the risk of  
concentration to dryness.

20

2. Method according to claim 1, characterised in that said  
force vector is sweeping the filter surface in a direction  
parallel to or with an angle away from at least part of the  
25 filter surface.

3. Method according to claim 1, characterised in that said  
force vector is sweeping the filter surface in a direction  
30 parallel to or with an angle into at least part of the filter  
surface.

4. Method according to claim 2, characterised in that the  
35 angle between said vector and the filter surface is comprised

in the interval 0 - 75°.

5. Method according to claim 3, characterised in that the  
5 angle between said vector and the filter surface is comprised  
in the interval 0 - 55°.

6. Method according to any of the claims 1 to 5,  
10 characterised in that the retentate in the concentrate pocket  
(4) is recovered by drawing the same from the pocket by means  
of a pipette or other recovery device.

15 7. Microconcentrator device for carrying out the method  
according to claim 1, comprising a sample reservoir (1) for  
receiving a sample to concentrate or purify, filter means (2)  
for filtering the sample, a centrifuge tube (5) for  
collecting the filtrate and a concentrate pocket (4) for  
20 retention of concentrate characterised in that said sample  
reservoir (1) is provided with a portion constituting a  
filtration chamber (3) with a volume being only a small  
fraction of the total volume of the sample reservoir having  
sidewalls and a bottom wall, that said filter means is  
25 arranged in the sidewall of said filtration chamber and that  
said concentrate pocket (4) is arranged in the bottom wall of  
said filtration chamber.

30 8. Microconcentrator device according to claim 7,  
characterised in that said filter means is a flat membrane  
provided with membrane supporting means mounted liquid tight  
in the side wall of said filtration chamber.



9. Microconcentrator device according to claim 7 or 8,  
characterised in that the external form of the filtration  
chamber (3) corresponds to half a cylinder, the filter means  
being arranged in the flat sidewall of said half cylinder, a  
5 corresponding half cylindrical sleeve (25) being arranged  
around said filtration chamber (3) keeping the filter means  
firmly fixed against the flat sidewall of said chamber (3).

10 10. Microconcentrator device according to claim 7 or 8,  
characterised in that the filtration chamber (3) has a  
tapered external form in the direction of the concentrate  
pocket (4), the filter means being arranged in a flat  
sidewall of said chamber (3), a sleeve (25) having a  
15 corresponding inner form being arranged around said  
filtration chamber (3) keeping the filter means firmly fixed  
against the flat sidewall.

20 11. Microconcentrator device according to claim 7,  
characterised in that said filter means is an outwardly  
convex membrane provided with membrane supporting means  
mounted liquid tight in the side wall of said filtration  
chamber.

25

12. Microconcentrator device according to claim 7,  
characterised in that said filter means is an outwardly  
concave membrane provided with membrane supporting means  
30 mounted liquid tight in the side wall of said filtration  
chamber.

13. Microconcentrator device according to any of the claims 7  
35 to 12, characterised in that said membrane has a maximum pore

size of 1 micrometer.

14. Microconcentrator device according to claim 7,  
5 characterised in that said filter means comprises a  
cylindrical porous membrane support constituting the side  
wall of the filtration chamber.

10 15. Microconcentrator device according to claim 14,  
characterised in that said cylindrical membrane support is  
mounted liquid tight and coaxially with the sample reservoir  
at the lower portion of said reservoir.

15  
16. Microconcentrator device according to claim 14 or 15,  
characterised in that said membrane support is made of a  
ceramic material.

20  
17. Microconcentrator device according to claim 14 or 15,  
characterised in that said membrane support is made of a  
carbon.

25  
18. Microconcentrator device according to claim 14,  
characterised in that the membrane is integrated on the  
surface of said membrane support.

30  
19. Microconcentrator device according to claim 14,  
characterised in that the membrane forms a separate  
cylindrical unit inserted in said membrane support.

35

20. Microconcentrator device according to any of the claims 7 to 19, characterised in that the bottom of the sample reservoir (1) is funnel-shaped with walls declining in the direction of the entry of the filtration channel, said walls  
5 being arranged with an angle between 5° and 46° to the direction perpendicular to the axis of the sample reservoir.

21. Microconcentrator device according to any of the claims 7  
10 to 20, characterised in that the sidewalls of the channel or the inside or outside of the surrounding sleeve is provided with indication markings making it possible to accurately indicate the remaining liquid volume.

15

22. Microconcentrator device according to any of the claims 7 to 21, characterised in that the dead stop feature is realized by means of a partition wall (28) separating the retentate reservoir (4) from the surface of the filter means  
20 and in that the cross section of the upper part of this wall has the shape of a ramp or a trapezoid.

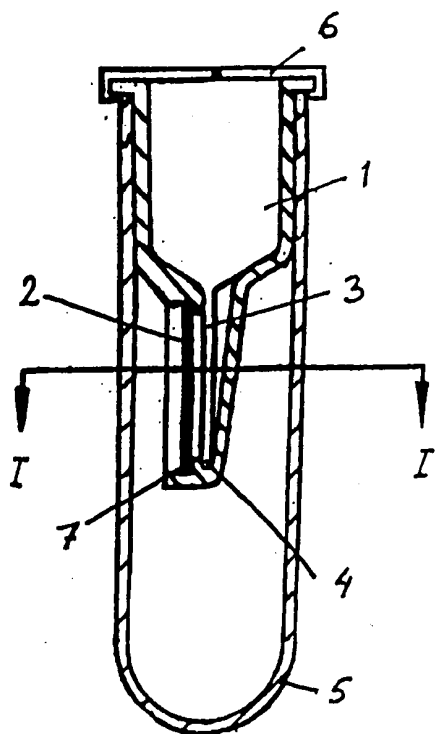


FIG 1

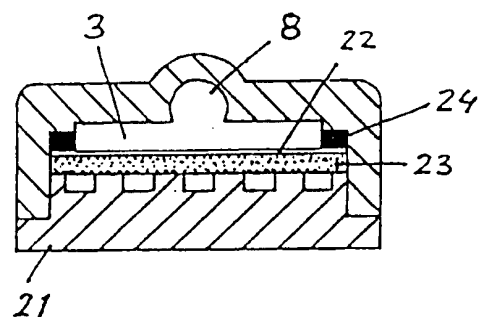


FIG 2

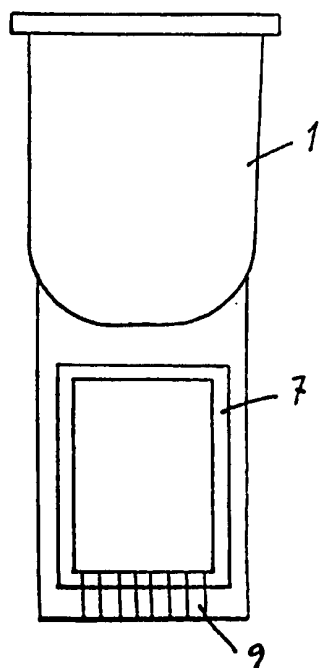


FIG 3

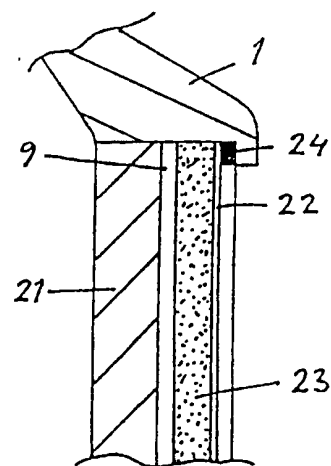


FIG 4

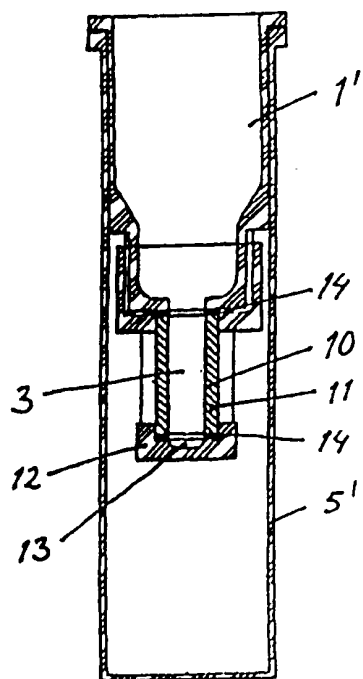


FIG 5

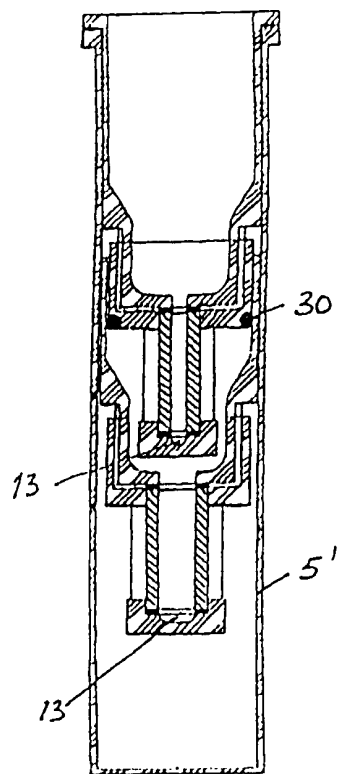


FIG 6

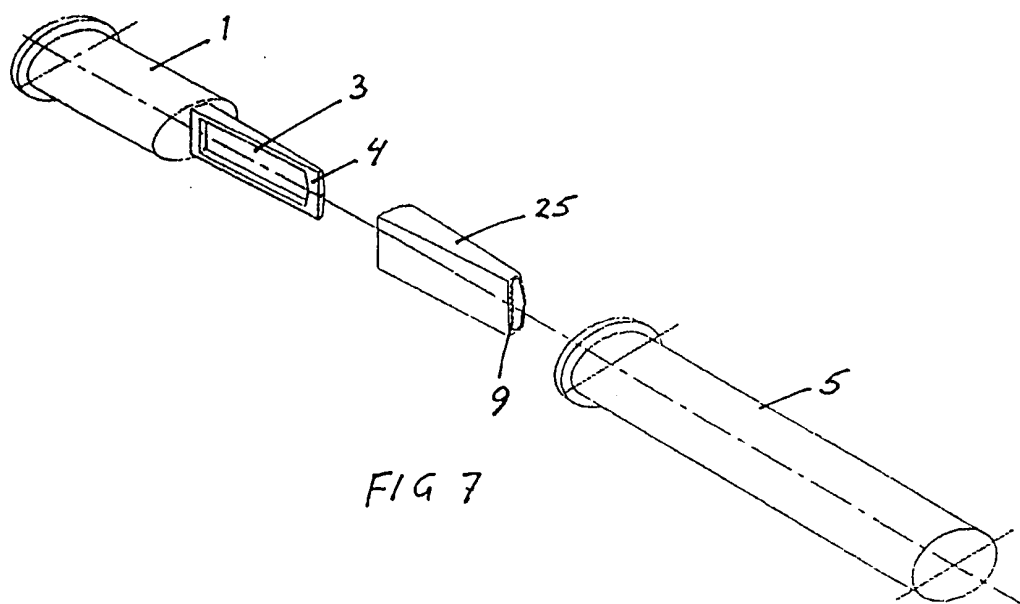


FIG 7

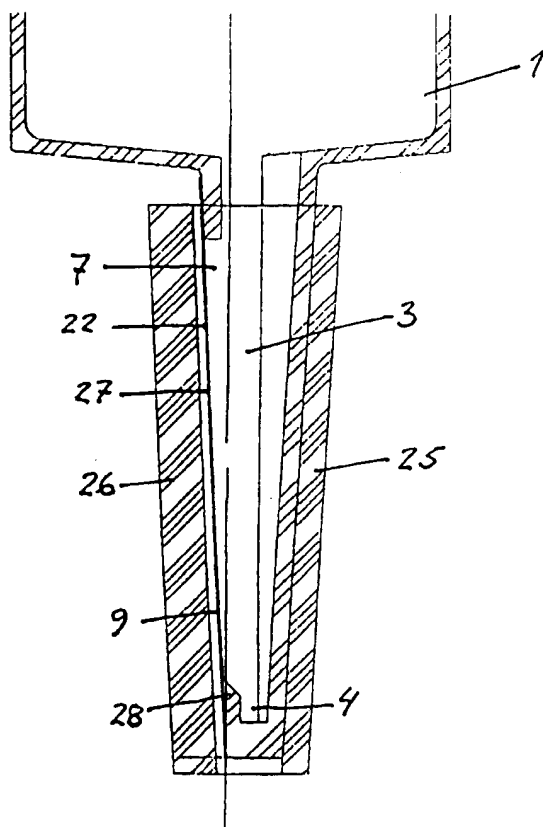


FIG 8

## INTERNATIONAL SEARCH REPORT

Internat. Application No

PCT/EP 94/01701

A. CLASSIFICATION OF SUBJECT MATTER  
 IPC 5 B01L3/14 B01D61/18 B01D63/16 B01D29/05 B01D29/74

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 5 B01L B01D

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US,A,3 488 768 (RIGOPOULOS) 6 January 1970 cited in the application see the whole document ---	1,7-15
A	US,A,4 632 761 (BOWERS ET AL.) 30 December 1986 cited in the application see the whole document ---	1
A	EP,A,0 430 355 (EASTMAN KODAK CO.) 5 June 1991 see the whole document ---	1-5
A	WO,A,89 00289 (SZABADOS) 12 January 1989 see the whole document ---	1
-/--		

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

## \* Special categories of cited documents:

- "A" document defining the general state of the art which is not considered to be of particular relevance
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Date of the actual completion of the international search

24 October 1994

Date of mailing of the international search report

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International Application No

PCT/EP 94/01701

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